

**PCT**

WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification <sup>6</sup> : <b>B01D 53/94</b>	A1	(11) International Publication Number: <b>WO 97/00119</b> (43) International Publication Date: <b>3 January 1997 (03.01.97)</b>
(21) International Application Number: <b>PCT/US96/07785</b>		(81) Designated States: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, UZ, VN, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).
(22) International Filing Date: <b>28 May 1996 (28.05.96)</b>		
(30) Priority Data: <b>08/490,892</b> <b>15 June 1995 (15.06.95)</b> <b>US</b>		
(71) Applicant: <b>ENGELHARD CORPORATION [US/US]</b> ; 101 Wood Avenue, Iselin, NJ 08830 (US).		
(72) Inventors: <b>DETTLING, Joseph, C.; 4 Spicy Pond Road, Howell, NJ 07731 (US). LUI, Yiu, Kwan; 3 Paprota Court, Parlin, NJ 08859 (US).</b>		Published <i>With international search report.</i>
(74) Agents: <b>MILLER, Stephen, I. et al.; Engelhard Corporation, 101 Wood Avenue, Iselin, NJ 08830 (US).</b>		

(54) Title: DIESEL ENGINE EXHAUST GAS CATALYST AND METHOD OF USE

(57) Abstract

A catalyst composition and structure containing the same and methods for treating diesel exhaust including a platinum group metal on a support in the presence of a catalyst activity controlling compound, a thermally stable ceria, and a non-catalytic pore-containing zeolite.

- 1 -

**Diesel engine exhaust gas catalyst and method of use.**

**BACKGROUND OF THE INVENTION**

**Field of the Invention**

5       The present invention relates to a catalyst composition and method of using the same for the oxidation of oxidizable components of exhaust gases, and more specifically to the treatment of diesel exhaust to reduce the content of particulates and other pollutants  
10       discharged to the atmosphere.

**Description of Related Art**

15       Diesel engine exhaust is a heterogeneous material which contains not only gaseous pollutants such as carbon monoxide ("CO") and unburned hydrocarbons ("HC"), but also soot particles which comprise both a dry, solid carbonaceous fraction and a soluble organic fraction. The soluble organic fraction is sometimes referred to as a volatile organic fraction ("VOF"), which terminology will be used herein. The VOF may exist in diesel exhaust  
20       either as a vapor or as an aerosol (fine droplets of liquid condensate) depending on the temperature of the diesel exhaust.

Oxidation catalysts comprising a platinum group metal

- 2 -

dispersed on a refractory metal oxide support are known for use in treating the exhaust of diesel engines in order to convert both HC and CO gaseous pollutants and particulates, i.e., soot particles, by catalyzing the 5 oxidation of these pollutants to carbon dioxide and water. One problem faced in the treatment of diesel engine exhaust is presented by the presence of sulfur in diesel fuel. Upon combustion, sulfur forms sulfur dioxide and the oxidation catalyst catalyzes the SO<sub>2</sub> to ,SO 10 ("sulfates") with subsequent formation of condensable sulfur compounds, such as sulfuric acid, which condense upon, and thereby add to, the mass of particulates. The sulfates also react with activated alumina supports to form aluminum sulfates, which render activated alumina- 15 containing catalysts inactive as disclosed in U.S. Patent 4,171,289. Previous attempts to deal with the sulfation problem include the incorporation of large amounts of sulfate-resistant materials such as vanadium oxide into the support coating, or the use of sulfur-resistant 20 support materials such as  $\alpha$ -alumina (alpha), silica and titania.

The prior art also shows an awareness of the use of zeolites, including metal-doped zeolites (i.e. catalytically active), to treat diesel exhaust. For 25 example, U.S. Patent 4,929,581 discloses a filter for diesel exhaust, in which the exhaust is constrained to

- 3 -

flow through the catalyst walls to filter the soot particles. A catalyst comprising a platinum group metal-doped zeolite is dispersed on the walls of the filter to catalyze oxidation of the soot to unplug the filter.

5 EPO 92/102161.4 discloses a catalyst for reducing the particle content and/or size in diesel engine exhaust by way of the zeolitic catalyst having acidic properties. The catalyst is stated to have properties enabling it to crack long-chain aromatic hydrocarbons. Zeolites include  
10 faujasite, pentasil and mordenite. Additionally, the faujasite and mordenite can be de-aluminated. The zeolite is stated to contain one or several transition elements which can include copper, nickel, cobalt, iron, chromium, manganese and/or vanadium.

15 Japanese Application No. S63-95026 (Publication No. H1-266854, October 24, 1989) discloses a catalyst for cleansing exhaust gas. The catalyst consists of zeolite, ion-exchanged with copper and carried on a fireproof carrier. The ion-exchange site is on the surface of the  
20 zeolite supercage and the coordination site of the oxygen atom for copper ion is a four-coordinate square.

Iwamoto, Catalytic Decomposition of Nitrogen Oxides, Petrotech 12, 888-893, 1989 is directed to the reduction of nitrogen oxides and various emissions from diesel  
25 exhaust. It is taught to use copper ion-exchanged ZSM-5 or mordanite or ferrierite. U.S. Patent No. 4,934,142

- 4 -

discloses an exhaust emission control device comprising a first filter provided in an exhaust system of an engine to collect particulates contained in an exhaust gas. A second filter is provided downstream of the first filter 5 to absorb an offensive odor component. The second filter is formed by an ion-exchange of copper ions of copper carried on a zeolite.

EPO Application No. 0 508 513 A1 discloses a method for treating diesel fuel engine exhaust to reduce emission 10 of particulates having cores of carbonaceous material and condensable hydrocarbons deposited on the carbonaceous material. The condensable hydrocarbons in the exhaust are contacted with a catalytically active solid acid material having hydrogen ions releasably retained at acidic sites 15 thereof. The condensable hydrocarbons in contact with the sites are cracked as hydrogen ions are released from the sites. The solid acid material is desirably a Y-type zeolite with (H) cations (HY zeolite), or hydrolyzed multi-valent cations such as lanthanum (LaY zeolite), 20 cerium (CeY zeolite) and calcium (CaY zeolite) and is supported on a ceramic or metal monolith.

U.S. Application No. 08/255,289 entitled, "Improved Zeolite-Containing Oxidation Catalyst and Method of Use" discloses a catalyst composition for treating a diesel 25 engine exhaust stream containing a volatile organic fraction. The catalyst composition comprises a refractory

- 5 -

carrier on which is disposed a coating of a catalytic material comprising a catalytically effective amount of ceria having a BET surface area of at least about 10 m<sup>2</sup>/g and a catalytically effective amount of a zeolite. It is 5 also known to employ an alumina stabilized ceria as a support for a platinum group metal as a dual exhaust catalyst.

The use of finely divided inorganic oxides containing vanadium and platinum group metal as active components is 10 disclosed in U.S. Patent No. 5,157,007. The catalyst is in the form of an open cell, monolith.

WO 94/22564 discloses a catalyst composition for treating diesel exhaust which includes ceria and optionally alumina as well as a beta zeolite. A platinum 15 group metal is employed to promote oxidation of CO and HC while limiting the conversion of SO<sub>2</sub> to SO<sub>3</sub>.

As is well-known in the art, catalysts used to treat the exhaust of internal combustion engines are less effective during periods of relatively low temperature 20 operation, such as the initial cold-start period of engine operation. This is because the engine exhaust is not at a temperature sufficiently high for the efficient catalytic conversion of noxious components in the exhaust. To this end, it is known in the art to employ high loads 25 of the platinum group metal catalyst to increase catalytic activity at low temperatures. It is also known to include

- 6 -

an adsorbent material, which may be a zeolite, as part of a catalytic treatment system in order to adsorb gaseous pollutants, usually hydrocarbons, and retain them during the initial cold-start period until the exhaust reaches a 5 more suitable, higher temperature. As the exhaust gas temperature increases, the adsorbed hydrocarbons are driven from the adsorbent and subjected to catalytic treatment at the higher temperature as disclosed, for example, in U.S. Patent 5,125,231 in which platinum group 10 metal-doped zeolites are employed as low temperature hydrocarbon adsorbents and oxidation catalysts.

Such efforts to improve upon the performance of diesel exhaust catalysts have been problematical. This is because, the low and high temperature operating 15 conditions, the presence of SO<sub>2</sub> and the need to effectively convert CO and HC to innocuous materials often impose competing requirements on diesel exhaust catalysts. For example, it is known that high loading of platinum group metals is required to convert CO and HC at low 20 temperatures. However, high loading of the platinum group metal increases the rate of conversion of SO<sub>2</sub> to SO<sub>3</sub>.

It is also known to modify the activity of the platinum group metals by adding appreciable amounts of vanadium oxide to the catalyst composition. Vanadium 25 oxide reduces the activity of the platinum metal to thereby reduce the rate at which SO<sub>2</sub> is converted to SO<sub>3</sub>.

- 7 -

However, after a relatively short operating period, vanadium oxide begins to irreversibly deactivate the platinum group metal thereby decreasing the performance of the catalyst in the conversion of CO and HC.

5 It would therefore be a significant advance in the art of converting diesel exhaust to innocuous materials to provide a catalyst which effectively converts CO and HC including the volatile organic fraction, while minimizing the conversion of SO<sub>2</sub> to SO<sub>3</sub>.

10

Summary of the Invention

The present invention is generally directed to a catalyst composition, structures containing the same and methods for oxidizing oxidizable components of a diesel engine exhaust stream in which at least some of a volatile 15 organic fraction of the diesel exhaust is converted to innocuous materials and in which gaseous hydrocarbons (HC) and carbon monoxide (CO) pollutants may also be similarly converted. The operation of the catalyst composition and the conversion reactions take place without the 20 substantial conversion of sulfur dioxide (SO<sub>2</sub>) to sulfur trioxide (SO<sub>3</sub>).

The catalyst composition for treating a diesel engine exhaust stream specifically in accordance with the present invention comprises:

25 a) a catalytically effective amount of at least one

- 8 -

platinum group metal on a support in the presence of at least one catalytic activity controlling compound;

- b) a thermally stable ceria; and
- c) a non-catalytic pore-containing zeolite.

5 In one aspect of the invention, the catalyst composition contains at least one compound which effectively controls the catalytic activity of the platinum group metal. The catalyst can therefore be employed in high loading amounts suitable for low  
10 temperature operation while minimizing the conversion of SO<sub>2</sub> to SO<sub>3</sub>. Preferred compounds for controlling catalytic activity are compounds containing vanadium, gold, silver and iron and combinations thereof.

The present catalyst composition employs a thermally  
15 stable ceria which effectively oxidizes the VOF (volatile organic fraction) of the diesel exhaust. The ceria component also serves to protect the platinum group metal from contact with the VOF to minimize the conversion of SO<sub>2</sub> to SO<sub>3</sub>, as well as to decrease coke formation.

20 In another aspect of the invention the catalyst composition employs a non-catalytic pore-containing zeolite which absorbs VOF at low temperatures and releases the same at higher temperatures, but generally below the temperature at which SO<sub>2</sub> converts to SO<sub>3</sub>. A particularly  
25 preferred zeolite is a hydrogen-beta zeolite.

The catalyst composition may be applied as a single

- 9 -

washcoat or as multiple coats (e.g. two coats), preferably with the thermally stable ceria as a top coat and the remaining components within a bottom coat. In this embodiment of the invention the ceria based top coat has 5 initial contact with the diesel exhaust stream to thereby absorb the VOF as well as to protect the platinum group metal.

Detailed Description of the Invention

As used herein and in the claims, the following terms 10 shall have the indicated meanings.

The term "washcoat" refers to a thin, adherent coating of a material, such as the catalytic material of the present invention, disposed on the walls forming the parallel gas flow passages of a carrier, which is 15 typically made of a refractory material such as cordierite or other oxide or oxide mixture, or a stainless steel.

The term "thermally stable ceria" means ceria that does not alter its physical structure at typical diesel exhaust gas temperatures of up to about 700°C.

20 The term "bulk form" for ceria means that the ceria is present as discrete particles (which may be, and usually are, of very small size, e.g., 10 to 20 microns in diameter or even smaller) as opposed to having been dispersed in solution form into another component.

25 Catalysts applied to diesel applications must deal

- 10 -

with many factors not associated with gasoline engines. Since the exhaust gas from diesel engines, especially those equipped with turbochargers, is cool at 150 - 200°C, it is imperative that the catalyst maintains activity at 5 very low temperatures. Fresh platinum metal catalysts oxidize CO and HC around 200°C in the presence of SO<sub>2</sub>. Until the temperature of the diesel exhaust stream reaches about 200°C, CO and HC are emitted into the atmosphere.

As previously described, diesel engine exhaust is 10 comprised of not only CO and unburnt HC but also a soot phase which includes a volatile organic fraction (VOF) including unburnt fuel and lubrication oil. The VOF, unless treated in advance, can deactivate the platinum metal catalyst until temperatures are reached sufficient 15 to oxidize these materials.

On the other hand, as temperatures of the exhaust stream rise, the SO<sub>2</sub> tends to oxidize to form SO<sub>3</sub> which negatively impacts total particulate matter (TPM) emissions. In particular, at temperatures above 300°C, the 20 rate at which platinum oxidizes SO<sub>2</sub> to SO<sub>3</sub> increases dramatically. Thus, in the operation of diesel engines deactivation of the catalyst can occur at lower temperatures while unwanted production of SO<sub>3</sub> can occur at higher temperatures. The catalyst composition of the 25 present invention addresses this problem by a) employing high loading amounts of the platinum group metal catalyst,

- 11 -

- b) controlling the activity of the platinum group metal,  
c) avoiding deactivation of the platinum group metal, d)  
storing HC's at lower temperatures in a non-catalytic  
environment, and e) minimizing the production of SO<sub>3</sub> when  
5 the exhaust stream reaches catalytically active  
temperatures.

The catalyst composition of the present invention employs three principal components, a platinum group metal component on a support including a catalytic activity  
10 controlling compound, a thermally stable ceria and an adsorbent, non-catalytic zeolite. The platinum group metal component as employed in the present invention is principally responsible for oxidizing gaseous HC and CO into innocuous materials such as water vapor and carbon  
15 dioxide without significantly catalyzing the conversion of SO<sub>2</sub> to SO<sub>3</sub>. The thermally stable ceria functions to oxidize liquid phase HC (VOF) while the zeolite component adsorbs gaseous HC at low, non-catalytic temperatures and then desorbs gaseous HC at temperatures generally below  
20 the temperature at which a significant amount of SO<sub>2</sub> converts to SO<sub>3</sub>. As a result, the platinum group metal converts HC to innocuous materials without converting a significant portion of adsorbed SO<sub>2</sub> to SO<sub>3</sub>.

The platinum group metal component of the present  
25 invention includes any and all platinum group metals alone or in combination including oxides thereof. The platinum

- 12 -

group metals include, for example, platinum, palladium, ruthenium, rhodium, iridium and mixtures and combinations of the same and their oxides. Platinum is the most preferred of the platinum group metals.

5       The amount of platinum group metal employed in the present catalyst composition should be a high loading quantity to maximize the conversion of CO and HC at the low initial temperatures of diesel fuel operation. The amount of the platinum group metal is generally at least  
10      about 5 g/ft<sup>3</sup>, typically in the range of from about 5 to 100 g/ft<sup>3</sup>, most preferably from about 10 to 70 g/ft<sup>3</sup>. The catalyst composition of the present invention therefore differs from low loading compositions in which the platinum group metal is used in much lower quantities,  
15      typically no more than about 2.0 g/ft<sup>3</sup>.

The support for the platinum group metal can be any support which does not tend to deactivate the platinum group metal during diesel exhaust treatment. Such supports include, zirconia, titania, silica and  
20      combinations thereof, preferably having a relatively low surface area. The preferred support is alumina, especially alumina having a relatively low surface area. The reduced surface area of the support serves to control the catalytic activity of the platinum group metal. While  
25      the surface area of the support may vary from about 50 to 200 m<sup>2</sup>/g, the preferred surface area is in the range of

- 13 -

from about 90 to 110 m<sup>2</sup>/g.

The platinum group metal component can be prepared, for example, in the manner taught in Saul G. Hinden, U.S. Patent No. 4,134,860, incorporated herein by reference.

5 A finely-divided, support (e.g. alumina) is contacted with a solution of a water-soluble, platinum group metal (e.g. platinum) to provide a composite which is essentially devoid of free or unabsorbed liquid. The platinum is converted into water-insoluble form while the composite  
10 remains essentially free of unabsorbed liquid. The composite is comminuted as a slurry to provide solid particles typically in the range of up to about 15 microns. The composite is then dried and calcined.

The catalytic component of the present invention is  
15 provided with an effective amount of at least one catalytic activity controlling material. This material serves to control (e.g. reduce) the catalytic activity of the platinum group metal so that high loadings of the platinum group metal can be employed for low temperature  
20 operations without a corresponding high conversion rate of adsorbed SO<sub>2</sub> to SO<sub>3</sub>.

While any material which can control the catalytic activity of the platinum group metal may be used, the preferred materials include compounds containing gold,  
25 vanadium, silver and iron and combinations thereof, such as oxides of gold, vanadium, silver and iron. The

- 14 -

starting materials for forming the catalytic activity controlling compounds are generally non-chloride, water-soluble compounds such as NaAuSO<sub>3</sub>, NH<sub>3</sub>VO<sub>3</sub>, V<sub>2</sub>O<sub>5</sub>, AgNO<sub>3</sub> and Fe(NO<sub>3</sub>)<sub>3</sub> · 9H<sub>2</sub>O, and the like. The amount of the catalytic 5 activity controlling material is typically in the range of from about 1 to 200 g/ft<sup>3</sup>, preferably from about 2 to 50 g/ft<sup>3</sup>.

Thermally stable ceria is employed in the present catalyst composition to adsorb SO<sub>2</sub> at low temperatures and 10 desorb the SO<sub>2</sub> below the temperature at which the platinum group metal vigorously catalyzes the conversion of adsorbed SO<sub>2</sub> to SO<sub>3</sub>. The ceria component also converts the VOF to innocuous materials. In particular, the ceria component must adsorb SO<sub>2</sub> under initial engine start up 15 conditions and desorb SO<sub>2</sub> at temperatures below about 300°C. In this way the SO<sub>2</sub> passes through the catalyst system under conditions which do not favor conversion to SO<sub>3</sub>. The amount of the ceria component of the catalyst is typically from about 10 to 60% by weight, preferably from 20 about 20 to 50% by weight, and most preferably from about 20 to 40% by weight, based on the total weight of the catalyst composition.

Ceria in bulk form is the preferred thermally stable ceria material for use in the catalyst composition. Bulk 25 ceria is solid, fine particulate ceria typically having a particle size distribution such that at least 95% by

- 15 -

weight of the particles have a diameter exceeding 0.5 microns. Further details regarding the structure and function of bulk ceria can be found in Chung-Zong Wan et al., U.S. Patent No. 4,714,694 incorporated herein by reference. It will be understood that the ceria component employed in the present catalyst should be thermally stable at temperatures of the diesel exhaust stream, typically up to 700°C.

The zeolite component of the catalyst composition adsorbs and retains gaseous HC at below catalytic temperatures. The zeolite does not itself catalyze any of the components of the diesel exhaust stream. Accordingly, the zeolite is non-catalytic and is not doped with catalytic materials such as platinum, iron and the like. The structure of the zeolite includes pores or cages which are capable of adsorbing and then desorbing HC. Desorption of the HC occurs when the diesel exhaust is at a high enough temperature to impart sufficient energy to the adsorbed HC molecules to enable them to escape the zeolite pores. Examples of the zeolite material meeting the criteria of the present invention include, for example, hydrogen-beta zeolite, Y-zeolite, pentasil, mordenite and mixtures thereof. Hydrogen-beta zeolite is the preferred zeolite.  $\beta$ -zeolites which may be employed in the present invention are described in Beck, Zeolite Molecular Sieves, Structure, Chemistry and Use, John Wiley

- 16 -

and Sons (1974); Bonetto et al., Optimization of Zeolite-Beta in Cracking Catalysts, Influence and Crystallite Size, Applied Catalysis, pp. 37-51 (1992); and U.S. Reissue Patent 28,341 of U.S. Patent No. 3,308,069; and  
5 Newsam et al., Structural Characterization of Zeolite Beta, Proc. R. Soc. Lond. A 420.375-405 (1988), each of which is incorporated herewith by reference.

The silica to alumina ratio for  $\beta$ -zeolite is from about 10 to about 200.  $\beta$ -zeolites are 12-member ring  
10 tridirectional zeolites with two types of channels, one being about 7.0 and the other about 5.5 angstroms. They are known to have larger pore sizes, high silica to alumina synthesis ratio and a tridirectional network of pores making them particularly suited for adsorbing HC.

15 The range of the amounts of the zeolite component is similar to that of ceria. Typically the zeolite component is present in an amount from about 10 to 60% by weight, preferably 20 to 50% by weight, most preferably 20 to 40% by weight, based on the total weight of the catalyst  
20 composition.

The carrier or substrate used in this invention should be relatively inert with respect to the catalytic composition dispersed thereon. The preferred carriers are comprised of ceramic-like materials such as cordierite,  $\alpha$ -alumina, silicon nitride, zirconia, mullite, spodumene, alumina-silica-magnesia, zirconium silicate, and

- 17 -

refractory metals such as stainless steel. The carriers are preferably of the type sometimes referred to as honeycomb or monolithic carriers, comprising a unitary body, usually cylindrical in configuration, having a plurality of fine, substantially parallel gas flow passages extending therethrough and connecting both end-faces of the carrier to provide a "flow-through" type of carrier. Such monolithic carriers may contain up to about 700 or more flow channels ("cells") per square inch of cross section, although far fewer may be used. For example, the carrier may have from about 7 to 600, more usually from about 200 to 400, cells per square inch ("cpsi").

Wall-flow carriers (filters) may also be used. Wall-flow carriers are generally similar in structure to flow-through carriers, with the distinction that each channel is blocked at one end of the carrier body, with alternate channels blocked at opposite end-faces. Wall-flow carrier substrates and the support coatings deposited thereon are necessarily porous, as the exhaust must pass through the walls of the carrier in order to exit the carrier structure.

The catalyst composition is deposited on the carrier such as a monolithic ceramic material in any conventional manner. A preferred method is to impregnate the carrier with an aqueous slurry of fine particles of the catalyst

- 18 -

composition. This can be accomplished by dipping the carrier (e.g. wall flow article) into the slurry, removing excess slurry by draining and subsequent drying at from about 100 to 150°C, followed by calcining at from about 450  
5 to 600°C.

The catalyst composition may also be applied to the carrier in multiple coats, typically as two coats. The composition of the respective coats will depend, in part, on the type of diesel exhaust being treated. For example,  
10 the support can be applied as a bottom coat and the platinum group metal, the catalytic activity controlling compound, the ceria and zeolite components as a top coat. The application of a dual coat may be conducted by first applying a slurry of the support onto the carrier followed  
15 by drying and calcining. The second coat is thereafter applied by first forming a slurry of the components of the second coat and applying the second coat in the same manner as the first coat.

In an embodiment of the invention particularly suited  
20 for diesel exhausts having a high VOF content, the bottom coat of the dual coat application contains the platinum group metal, the catalytic activity controlling compound, the support and the zeolite while the top coat contains the thermally stable ceria. By employing ceria as the top  
25 coat, there is a more effective oxidation of VOF and better protection of the platinum group metal from the

- 19 -

e deactivating effects of contact with VOF.

g In another embodiment of the catalyst composition of  
n the present invention which is particularly effective when  
0 the diesel exhaust has a dry soot content, the bottom coat  
e 5 includes the ceria and zeolite components while the top  
e coat contains the platinum group metal, catalytic activity  
controlling compound and the support. In other  
embodiments of the invention, at least one of the ceria  
and zeolite components are provided in each of the coats  
e 10 of the catalyst composition.

#### EXAMPLE 1

A catalyst composition according to the present invention was formed by preparing a first material starting with an ammoniacal solution containing 80% by weight of the total platinum metal employed in the catalyst composition and combining the same with 396 g of alumina having a surface area of about 90 m<sup>2</sup>/g. The platinum and alumina were pre-mixed followed by the addition of acetic acid in an amount of 4% by weight based 15 on the weight of the alumina. After mixing, a solution of ferric nitrate [Fe(NO<sub>3</sub>)<sub>3</sub> · 9H<sub>2</sub>O], providing a Pt/Fe weight ratio of 10 was added and the combined solution mixed and ball milled in the presence of added water to give a mixture having a 46% solids content.  
20

25 A second material was prepared by starting with an

- 20 -

ammoniacal solution containing 20% by weight of the total platinum metal employed in the catalyst composition. The platinum solution was combined with bulk ceria in an amount of 396 g and premixed to form the second material.

5       The first and second materials as well as 396 g of H-beta zeolite were blended in the presence of water to provide a washcoat slurry having a 49% solids content. The slurry was milled to a mean particle size of less than 8 $\mu$ . The resulting slurry, having a solids content of 48 -  
10      49%, a pH of 3.5 to 3.8 and a viscosity of 20 to 30 cps, was coated on a monolithic cordierite substrate (400 cells/in<sup>2</sup>) in an amount sufficient to provide a washcoat gain of 1.95 g/in<sup>3</sup> and then dried at 100 to 150°C and calcined at about 450°C. The final catalyst composition  
15      contained 20 g/ft<sup>3</sup> of platinum and 0.8 g/ft<sup>3</sup> of iron.

#### EXAMPLE 2

A slurry containing 600 g of alumina having a surface area of 150 m<sup>2</sup>/g, 200 g of bulk ceria and 200 g of H-beta zeolite were blended together to give a mixture having a  
20      solids content of 36%. The mixture was milled so that 90% of the particles were less than 8 $\mu$ . The slurry was coated on a monolithic cordierite substrate (400 cells/in<sup>2</sup>) at a washcoat gain of 1.0 g/in<sup>3</sup>, dried at about 100°C and calcined at 450°C to form a bottom catalyst layer.

25      A slurry containing platinum metal and vanadium oxide

- 21 -

from an ammoniacal solution was combined with 50 m<sup>2</sup>/g titania with the subsequent addition of 15 ml of acetic acid and then milled. The resulting slurry was combined with 300 g of bulk ceria, 400 g of H-beta zeolite and 100 5 g of SiO<sub>2</sub> from SiO<sub>2</sub> sol solution and blended together to form a slurry with 90% of the particles having a particle size of less than 8μ. The slurry was coated onto the above-formed bottom catalyst layer to form a top catalyst layer having a washcoat gain of 1.5 g/in<sup>3</sup>. The coated 10 substrate was dried at 105°C and calcined at 450°C to provide a catalyst containing 40 g/ft<sup>3</sup> of platinum and 30 g/ft<sup>3</sup> of vanadium.

EXAMPLE 3

A bottom catalyst layer having the same composition 15 as described in Example 2 was applied to a monolithic substrate made of cordierite.

A top catalyst layer slurry was prepared. 427 g of a 25% by weight ZrO<sub>2</sub>/SiO<sub>2</sub> composite (210 m<sup>2</sup>/g) material was placed in a vessel. An ammoniacal solution of platinum as 20 employed in Example 1 (14.12 g of platinum) and 0.70 g of gold from a NaAuSO<sub>4</sub> solution were placed in a vessel and diluted to 450 ml with deionized water. The combined platinum-gold solution was added to the ZrO<sub>2</sub>/SiO<sub>2</sub> composite material and mixed to obtain a uniform mixture.

25 15 ml of acetic acid was slowly added to the uniform

- 22 -

mixture followed by 20 ml of formic acid under continuous mixing. The mixed solution was transferred to a ball mill to which was added 183 g of thermally stable ceria, 244 g of H-beta zeolite, 203 g of SiO<sub>2</sub> sol solution (30% SiO<sub>2</sub>) 5 and 925 g of deionized water. The solution was milled until 90% of the particles had a particle size of less than 8μ.

The slurry was coated on the bottom catalyst layer at a washcoat gain of 1.5 g/in<sup>3</sup>. The substrate was dried at 10 100°C and calcined at 450°C. The resulting catalyst contained 40 g/ft<sup>3</sup> of platinum and 2 g/ft<sup>3</sup> of gold.

#### EXAMPLE 4

A slurry containing 700 g of alumina having a surface area of 90 m<sup>2</sup>/g and 300 g of bulk ceria were blended with 15 water to give a mixture having a solids content of 38%. The mixture was milled until 90% of the particles had a particle size of less than 8μ and then coated on a monolithic cordierite substrate (400 cells/in<sup>2</sup>) at a washcoat gain of 1.0 g/in<sup>3</sup>, dried at 105°C and calcined at 20 450°C to form a bottom catalyst layer.

A mixture of an ammoniacal solution of platinum, a solution of Pd(NH<sub>3</sub>)<sub>4</sub>(NO<sub>3</sub>)<sub>2</sub> and a solution of NaAuSO<sub>3</sub> were combined with 500 g of a ZrO<sub>2</sub>/SiO<sub>2</sub> composite and mixed followed by the addition of acetic acid and formic acid.

25 The resulting slurry was combined with 300 g of bulk

- 23 -

ceria, 600 g of H-beta zeolite, 50 g of  $\text{Pr}_2\text{O}_3$  and a sufficient amount of water to give a slurry with a 37% solids content. The slurry was milled until 90% of the particles had a particle size of less than 8 microns. The 5 slurry was applied to the bottom catalyst layer at a washcoat gain of 1.45 g/in<sup>3</sup>, dried at about 105°C and calcined at about 500°C. The resulting catalyst contained 40 g/ft<sup>3</sup> of platinum, 1 g/ft<sup>3</sup> of palladium and 2 g/ft<sup>3</sup> of gold.

10

EXAMPLE 5

The same bottom catalyst layer described in Example 4 was formed on a monolithic substrate. A top catalyst layer was formed as described in Example 4 except that the amount of gold was increased to 5 g/ft<sup>3</sup>.

15

EXAMPLE 6

500 g of a 12%  $\text{TiO}_2\text{-Al}_2\text{O}_3$  composite material was mixed with an ammoniacal solution (diluted to 400 ml with deionized water) containing 14.47 g of platinum. To this mixture was added 15 ml of acetic acid under mixing.

20

A solution of  $\text{AgNO}_3$  containing 1.8 g of Ag was diluted to 25 ml with deionized water. The diluted silver solution was added to the  $\text{Pt-TiO}_2\text{-Al}_2\text{O}_3$  produced above and mixed.

The mixture was combined with 250 g of H-beta zeolite

- 24 -

and 475 g of deionized water. The resulting slurry was milled until 90% of the particles had a particle size of less than 8 $\mu$ .

A cordierite honeycomb substrate (400 cells/in<sup>2</sup>) was  
5 coated with the slurry at a washcoat gain of 1.2 g/in<sup>3</sup>. The coated substrate was dried at 100°C and calcined at 450°C to form a bottom catalyst layer.

A top catalyst layer, prepared in the following manner, was coated on the bottom catalyst layer.

10 600 g of gamma-alumina, 400 g of bulk ceria, and 1500 ml of deionized water were combined and milled until 90% of the particles had a particle size of less than 8 $\mu$ . The slurry was then coated on the bottom catalyst layer at a washcoat gain of 1.0 g/in<sup>3</sup>. The top coating layer was then  
15 dried at 100°C and calcined at 450°C. The resulting catalyst contained 40 g/ft<sup>3</sup> of platinum and 5 g/ft of silver.

#### EXAMPLE 7

438 g of a 25% ZrO<sub>2</sub>-SiO<sub>2</sub> composite material was mixed  
20 with an ammoniacal solution (diluted to 350 ml with deionized water) containing 14.47 g of platinum. To this mixture was added 15 ml acetic acid under mixing.

A solution of NaAuSO<sub>4</sub> containing 1.81 g Au was diluted to 25 ml with deionized water. The diluted gold solution

- 25 -

was added to the Pt-ZrO<sub>2</sub>-SiO<sub>2</sub> produced above and mixed.

The mixture was combined with 250 g of H-beta zeolite, 250 g of bulk ceria, 310 g of a 30% solution of SiO<sub>2</sub> and 500 g of deionized water. The resulting slurry  
5 was milled until 90% of the particles had a particle size of less than 8μ.

A cordierite honeycomb substrate (400 cells/in<sup>2</sup>) was coated with the slurry at a washcoat gain of 1.6 g/in<sup>3</sup>. The coated substrate was dried at 100°C and calcined at  
10 450°C to form a bottom catalyst layer.

A top catalyst layer prepared in the following manner was coated on the bottom catalyst layer.

400 g of a 2% SiO<sub>2</sub> doped titania, 400 g of bulk ceria, 100 g of H-beta zeolite, 500 g of a 30% solution of SiO<sub>2</sub>  
15 and 1100 ml of deionized water were combined and milled until 90% of the particles had a particle size of less than 8μ. The slurry was then coated on the bottom catalyst layer at a washcoat gain of 1.0 g/in<sup>3</sup>. The top catalyst layer was then dried at 100°C and calcined at  
20 450°C. The resulting catalyst contained 40 g/ft<sup>3</sup> of platinum and 5 g/ft<sup>3</sup> of gold.

#### REFERENCE EXAMPLE 1

A reference catalyst composition (Ref. Ex. 1) was prepared in the following manner.

25 396 g of gamma-alumina was placed in a vessel and

- 26 -

mixed. An ammoniacal solution of platinum as employed in Example 1 (6.336 g of platinum) was placed in a separate vessel and diluted to 270 ml with deionized water. The platinum solution was slowly added to the alumina and 5 mixed followed by the addition of 15 ml of concentrated acetic acid and further mixing. The resulting solution and 450 g of deionized water were placed in a ball mill and milled until 90% of the particles had a particle size of less than 8 $\mu$ .

10 A slurry containing platinum/ceria and H-beta zeolite was prepared in the following manner.

396 g of ceria-zirconia composite material was placed in a vessel and mixed. An ammoniacal solution of platinum as used in Example 1 (0.704 g of platinum) was placed in 15 a separate vessel and diluted to 120 ml with deionized water. The platinum solution was added to the solution of ceria-zirconia and mixed until uniform. 12 ml of concentrated acetic acid was added to the uniform solution followed by mixing. The resulting solution as well as 396 20 g of H-beta zeolite and 650 g of deionized water were placed in a ball mill and milled until 90% of the particles had a particle size of less than 8 $\mu$ .

The two slurries were combined and blended and coated on a cordierite honeycomb substrate (400 cells/in<sup>2</sup>) at a 25 washcoat gain of 1.95 g/in<sup>3</sup>. The substrate was dried at 100°C and calcined at 450°C. The resulting catalyst

- 27 -

contained 20 g/ft<sup>3</sup> of platinum.

Reference Example 1 and the catalyst composition prepared in accordance with Example 1 were used to treat a diesel exhaust stream containing C<sub>7</sub>H<sub>16</sub>, C<sub>3</sub>H<sub>8</sub>, CO, SO<sub>2</sub> and 5 H<sub>2</sub>O under the following conditions.

A reactor synthetic gas having the following composition

200 ppm HC as propylene and propane in  
a 2:1 ratio

10 200 ppm CO

1000 ppm NO

50 ppm SO<sub>2</sub>

10% by volume H<sub>2</sub>O (steam)

4.5% by volume CO<sub>2</sub>

15 10% by volume O<sub>2</sub>

balance N<sub>2</sub>

was provided at a space velocity of 50,000 (volume hour, 1/hr) at an aging temperature of 500°C for 2 hours in reactor gases. The size of the catalyst was 87 cm<sup>3</sup> with 62 20 cells/cm<sup>2</sup>.

Each of the catalyst compositions were tested over a range of operating temperatures of from 250°C to 400°C in increments of 50°C. The % conversion of HC, CO and SO<sub>2</sub> were measured by individual analyzers and the results 25 shown in Table 1.

- 28 -

**TABLE 1**

Temperature	Ref. Example 1 % Conversion			Example 1 % Conversion		
	HC	CO	SO <sub>2</sub>	HC	CO	SO <sub>2</sub>
5	250°C	62	98	27	59	96
	300°C	67	98	31	63	98
	350°C	78	98	60	70	98
	400°C	81	98	72	75	98

As shown in Table 1 the % conversion of SO<sub>2</sub> to SO<sub>3</sub> was significantly lower for the catalyst of Example 1 as compared with Reference Example 1, particularly at temperatures of 300°C or more. This shows that the catalytic activity of the platinum metal is effectively controlled in the present invention to limit the undesirable formation of sulfates. The present catalyst also provides sufficient conversion rates of HC and CO comparable to that of the reference catalyst.

#### REFERENCE EXAMPLE 2

A reference catalyst composition (Ref. Ex. 2) was prepared in the following manner.

420 g of gamma-alumina was placed in a vessel and mixed. An ammoniacal solution of platinum as employed in Example 1 (5.79 g of platinum) was placed in a separate vessel and diluted to 300 ml with deionized water. The

- 29 -

platinum solution was slowly added to the alumina and mixed followed by the addition of 12 ml of concentration acetic acid and further mixing. The resulting solution and 300 g of deionized water were placed in a ball mill 5 and milled until 90% of the particles had a particle size of less than 8 $\mu$ .

A slurry containing platinum/ceria and Fe-beta zeolite was prepared in the following manner.

415 g of alumina doped ceria was placed in a vessel 10 and mixed. An ammoniacal solution of platinum as used in Example 1 (5.79 g of platinum) was placed in a separate vessel and diluted to 125 ml with deionized water. The platinum solution was added to the solution of alumina doped ceria and mixed until uniform. 12 ml of 15 concentrated acetic acid was added to the uniform solution followed by mixing. The resulting solution as well as 415 g of Fe-beta zeolite and 700 g of deionized water were placed in a ball mill and milled until 90% of the particles had a particle size of less than 8 $\mu$ .

20 The two slurries were combined and blended and coated on a cordierite honeycomb substrate at a washcoat gain of 2.50 g/in<sup>3</sup>. The substrate was dried at 100°C and calcined at 450°C. The resulting catalyst contained 40 g/ft<sup>3</sup> of platinum.

25 Reference Example 2 and each of Examples 2 - 4 were used to treat a diesel exhaust stream containing C<sub>7</sub>H<sub>16</sub>, CO,

- 30 -

SO<sub>2</sub> and H<sub>2</sub>O under the same reactor conditions described previously for the comparison of Reference Example 1 and Example 1. Each of the catalyst compositions were tested over a range of operating temperatures in 50°C increments of from 200°C to 400°C. The % conversion at 50°C increments of HC, CO and SO<sub>2</sub> were measured and the results shown in Table 2.

**TABLE 2**

Temp °C	Reference Example 2 % Conversion			Example 2 % Conversion			Example 3 % Conversion			Example 4 % Conversion		
	HC	CO	SO <sub>2</sub>	HC	CO	SO <sub>2</sub>	HC	CO	SO <sub>2</sub>	HC	CO	SO <sub>2</sub>
200	4	98	33	4	94	12	2	98	28	3	97	6
250	46	98	55	23	97	16	28	98	28	27	97	16
300	92	98	55	82	97	20	87	98	41	87	98	46
350	96	99	68	96	98	29	96	98	50	94	98	66
400	97	99	76	97	98	49	97	98	63	95	98	76

As shown in Table 2, the % conversion of SO<sub>2</sub> to SO<sub>3</sub> for the present catalyst composition is significantly less than the reference catalyst especially at low operating temperatures.

Reference Example 1 and each of Examples 5 and 6 were used to treat the same diesel exhaust stream under the same reactor conditions as previously described above for Examples 2 - 4. Each of the catalyst compositions were measured for HC, CO and SO<sub>2</sub> conversion rates in the same

- 31 -

manner as described above and the results are shown in Table 3.

**TABLE 3**

Temp. °C	Reference Example 2 % Conversion			Example 5 % Conversion			Example 6 % Conversion		
	HC	CO	SO <sub>2</sub>	HC	CO	SO <sub>2</sub>	HC	CO	SO <sub>2</sub>
200	4	98	33		2	91	12		1
250	46	98	55		19	97	15		3
300	92	98	55		73	97	33		31
350	96	99	68		91	97	53		85
400	97	99	76		93	97	67		93
									27

As shown in Table 3, the % conversion of SO<sub>2</sub> to SO<sub>3</sub> for the present catalyst composition is significantly less than the reference catalyst.

Reference catalyst 2 and the catalyst of Example 7 were subjected to a diesel engine light-off activity test in the following manner.

A cordierite honeycomb substrate having a volume of 55 in<sup>3</sup>, and 400 cells/in<sup>2</sup>, was loaded with the respective catalysts having a platinum loading of 40 g/ft<sup>3</sup>. The catalyst structures were contacted with a diesel exhaust stream from a diesel engine operating at a speed effective

- 32 -

to provide 2000 l/min of exhaust, an engine load of 15 - 180 NM and a catalyst inlet temperature ranging from 100 to 530°C. The engine aging cycle was 130°C for 15 minutes followed by 650°C for 15 minutes.

- 5 The percentage conversion of HC, CO and total particulate matter (TPM) was measured and the results are shown in Table 4.

**TABLE 4**

Temp. °C	% HC Conv.		% CO Conv.		% TPM Conv.	
	Ref. 2	Ex. 7	Ref. 2	Ex. 7	Ref. 2	Ex. 7
150	58	65	7	5	45	59
200	41	92	11	5	66	71
250	57	70	86	78	50	55
300	79	82	86	95	40	59
350	72	76	72	94	11	19
400	71	65	76	91	5	16
450	72	65	75	92	-10	18
500	80	57	74	93	-30	25

- 33 -

As shown in Table 4, the conversion of TPM for the present catalyst composition was significantly greater than for the reference catalyst, especially at high operating temperatures while the present catalyst 5 exhibited sufficient conversion rates of HC and CO although somewhat less than the reference catalyst.

What is claimed is:

1. A catalyst composition for treating a diesel engine exhaust stream comprising:
  - a) a catalytically effective amount of at least one platinum group metal on a support in the presence of at least one catalyst activity controlling compound;
  - b) a thermally stable ceria; and
  - c) non-catalytic pore-containing zeolite.
- 10 2. The catalyst composition of claim 1 wherein the catalyst activity controlling compound is selected from compounds containing vanadium, gold, silver, iron and combinations thereof.
- 15 3. The catalyst composition of claim 1 wherein the amount of the catalyst activity controlling compound is from about 1 to 200 g/ft<sup>3</sup>.
4. The catalyst composition of claim 1 wherein the amount of the catalyst activity controlling compounds is from about 2 to 50 g/ft<sup>3</sup>.
- 20 5. The catalyst composition of claim 1 wherein the amount of the platinum group metal is at least about 5

- 35 -

g/ft<sup>3</sup>.

6. The catalyst composition of claim 1 wherein the amount of the platinum group metal is from about 5 to 100 g/ft<sup>3</sup>.

5 7. The catalyst composition of claim 1 wherein the amount of the platinum group metal is from about 10 to 70 g/ft<sup>3</sup>.

8. The catalyst composition of claim 1 wherein the support for the platinum group metal is selected from the  
10 group consisting of alumina, zirconia, titania, silica and combinations thereof.

9. The catalyst composition of claim 1 wherein the support for the platinum group metal is alumina.

10. The catalyst composition of claim 1 wherein the  
15 support has a low surface area.

11. The catalyst composition of claim 10 wherein the surface area of the support is from about 50 to 200 m<sup>2</sup>/g.

12. The catalyst composition of claim 10 wherein the

- 36 -

surface area of the support is from about 90 to 110 m<sup>2</sup>/g.

13. The catalyst composition of claim 1 wherein the thermally stable ceria comprises bulk form ceria composed of fine particles wherein 95% by weight of the particles 5 having a diameter exceeding 0.5μ.

14. The catalyst composition of claim 1 wherein the amount of the thermally stable ceria is from about 10 to 60% by weight based on the total weight of the catalyst composition.

10 15. The catalyst composition of claim 1 wherein the zeolite is selected from the group consisting of hydrogen-beta zeolite, Y-zeolite, pentasil, mordenite and mixtures thereof.

15 16. The catalyst composition of claim 1 wherein the zeolite is hydrogen-beta zeolite.

17. The catalyst composition of claim 1 wherein the amount of the zeolite is from about 10 to 60% by weight based on the total weight of the catalyst composition.

20 18. The catalyst composition of claim 1 wherein the platinum group metal is platinum.

- 37 -

19. A catalyst structure comprising:

- a) a catalyst receiving substrate; and
- b) the catalyst composition of claim 1 on said substrate.

5 20. The catalyst structure of claim 19 wherein the substrate is in the form of a flow-through carrier.

21. The catalyst structure of claim 19 wherein the substrate is in the form of a wall-flow carrier.

10 22. The catalyst structure of claim 19 wherein the catalyst composition is in the form of at least one washcoat.

15 23. The catalyst structure of claim 19 wherein the catalyst composition is in the form of two washcoats, a bottom washcoat comprising the support and a top washcoat comprising the platinum group metal, the catalytic activity controlling compound, the thermally stable ceria and the non-catalytic pore-containing zeolite.

20 24. The catalyst structure of claim 19 wherein the catalyst composition is in the form of two washcoats, a bottom washcoat comprising the platinum group metal, the catalytic activity controlling compound, the support and

- 38 -

the non-catalytic pore-containing zeolite and a top washcoat comprising the thermally stable ceria.

25. The catalyst structure of claim 19 wherein the catalyst composition is in the form of two washcoats, a  
5 bottom washcoat comprising the thermally stable ceria and the non-catalytic pore-containing zeolite and a top washcoat comprising the platinum group metal, the catalytic activity controlling compound and the support.

26. The catalyst structure of claim 19 wherein the catalyst composition is in the form of two washcoats, each  
10 coat containing at least one of said thermally stable ceria and non-catalytic pore-containing zeolite.

27. A method of treating a diesel exhaust stream comprising passing said diesel exhaust stream into  
15 operative contact with the catalyst composition of claim 1.

28. A method of treating a diesel exhaust stream comprising passing said diesel exhaust stream into operative contact with the catalyst structure of claim 19.

# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/US 96/07785

**A. CLASSIFICATION OF SUBJECT MATTER**  
IPC 6 B01D53/94

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)  
IPC 6 B01D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 94 22564 A (ENGELHARD CORP) 13 October 1994 see page 9, line 3 - page 14, line 16; claims 1-38 ---	1-12, 14-28
X	US 5 157 007 A (DOMESLE RAINER ET AL) 20 October 1992 see claims 1-22 ---	1-9,14, 18-22, 27,28
X	EP 0 559 021 A (DEGUSSA) 8 September 1993 see page 3, line 7 - line 37; claims 1-10; example 11; table 2 ---	1-9, 14-22, 26-28 -/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

\* Special categories of cited documents :

- \*'A' document defining the general state of the art which is not considered to be of particular relevance
- \*'E' earlier document but published on or after the international filing date
- \*'L' document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*'O' document referring to an oral disclosure, use, exhibition or other means
- \*'P' document published prior to the international filing date but later than the priority date claimed

\*'T' later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

\*'X' document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

\*'Y' document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

\*'&' document member of the same patent family

1

Date of the actual completion of the international search	Date of mailing of the international search report
18 September 1996	27.09.96
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Authorized officer  Eijkenboom, A

## INTERNATIONAL SEARCH REPORT

International Application No  
PCT/US 96/07785

## C(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5 407 880 A (IKEDA TAKUYA ET AL) 18 April 1995 see the whole document ---	1,3-9, 15-26
A	US 4 760 044 A (JOY III GEORGE C ET AL) 26 July 1988 see claims 1-31 ---	1-26
A	US 4 714 694 A (WAN CHUNG-ZONG ET AL) 22 December 1987 see column 8, line 16 - line 46; claims 1-52 -----	1-14, 18-22

**INTERNATIONAL SEARCH REPORT**

Information on patent family members

International Application No	
PCT/US 96/07785	

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO-A-9422564	13-10-94	CA-A-	2159317	13-10-94
		EP-A-	0691883	17-01-96
-----				
US-A-5157007	20-10-92	DE-A-	3940758	13-06-91
		AU-B-	629433	01-10-92
		AU-A-	6769990	13-06-91
		CA-A-	2031762	10-06-91
		CN-A-	1052262	19-06-91
		DE-D-	59005194	05-05-94
		EP-A-	0432534	19-06-91
		ES-T-	2023787	01-07-94
		JP-A-	3224631	03-10-91
		RU-C-	2022643	15-11-94
		TR-A-	25872	01-09-93
		US-A-	5514354	07-05-96
-----				
EP-A-0559021	08-09-93	DE-A-	4206699	09-09-93
		AT-T-	129427	15-11-95
		AU-A-	3395393	09-09-93
		BR-A-	9300731	08-09-93
		CA-A-	2090882	05-09-93
		CN-A-	1079413	15-12-93
		CZ-A-	9203799	16-02-94
		DE-D-	59300801	30-11-95
		ES-T-	2079219	01-01-96
		HU-A,B	66374	28-11-94
		JP-A-	6071179	15-03-94
		PL-A-	297926	15-11-93
		TR-A-	26500	15-03-95
		US-A-	5354720	11-10-94
		ZA-A-	9301486	04-10-93
-----				
US-A-5407880	18-04-95	JP-A-	6142519	24-05-94
		JP-A-	6198164	19-07-94
-----				
US-A-4760044	26-07-88	WO-A-	9000441	25-01-90
		CA-A-	1317274	04-05-93
		DE-T-	3870256	21-05-92
		US-A-	5008090	16-04-91
		AU-B-	619828	06-02-92

## INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No  
PCT/US 96/07785

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US-A-4760044		AU-A- 3350689 EP-A- 0425488	05-02-90 08-05-91
US-A-4714694	22-12-87	EP-A- 0251752 JP-A- 63007841 KR-B- 9502218	07-01-88 13-01-88 15-03-95

THIS PAGE BLANK (USPTO)